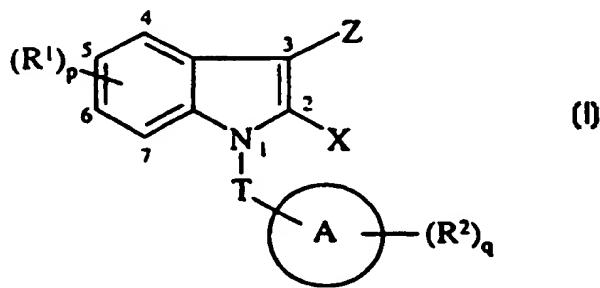




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(71) Applicant (for all designated States except US): ZENECA LIMITED [GB/GB]; 15 Stanhope Gate, London W1Y 6LN (GB).			
(72) Inventors; and			Published
(75) Inventors/Applicants (for US only): BARKER, Andrew, John [GB/GB]; Alderley Park, Macclesfield, Cheshire SK10 4TG (GB). KETTLE, Jason, Grant [GB/GB]; Alderley Park, Macclesfield, Cheshire SK10 4TG (GB). FAULL, Alan, Wellington [GB/GB]; Alderley Park, Macclesfield, Cheshire SK10 4TG (GB).			With international search report.
(74) Agent: DENERLEY, Paul, Millington; Intellectual Property Dept., Zeneca Pharmaceuticals, Mereside, Alderley Park, Macclesfield, Cheshire SK10 4TG (GB).			

(54) Title: INDOLE DERIVATIVES AS MCP-1 RECEPTOR ANTAGONISTS



(57) Abstract

The invention concerns pharmaceutically useful indoles of formula (I), in which Z, X, T, A, R¹, R², p and q have any of the meanings defined herein, and their pharmaceutically acceptable salts or *in vivo* hydrolysable esters, as well as pharmaceutical compositions containing them. The novel compounds possess inhibitory activity against monocyte chemoattractant protein-1 (MCP-1). The invention further concerns the use of such indoles in the treatment of a disease or condition mediated by MCP-1.

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INDOLE DERIVATIVES AS MCP-1 RECEPTOR ANTAGONISTS

The present invention relates to anti-inflammatory compounds that act via inhibition of Monocyte Chemoattractant Protein-1 (MCP-1) and in particular MCP-1 inhibitor compounds that contain an indole moiety. The invention further relates to processes for their preparation, 5 to intermediates useful in their preparation, to their use as therapeutic agents and to pharmaceutical compositions containing them.

MCP-1 is a member of the chemokine family of pro-inflammatory cytokines which mediate leukocyte chemotaxis and activation. MCP-1 is a C-C chemokine which is one of the most potent and selective T-cell and monocyte chemoattractant and activating agents known. 10 MCP-1 has been implicated in the pathophysiology of a large number of inflammatory diseases including rheumatoid arthritis, glomerular nephritis, lung fibrosis, restenosis (International Patent Application WO 94/09128), alveolitis (Jones et al., 1992, *J. Immunol.*, 149, 2147) and asthma. Other disease areas where MCP-1 is thought to play a part in their pathology are atherosclerosis (e.g. Koch et al., 1992, *J. Clin. Invest.*, 90, 772 -779), psoriasis 15 (Deleuran et al., 1996, *J. Dermatological Science*, 13, 228-236), delayed-type hypersensitivity reactions of the skin, inflammatory bowel disease (Grimm et al., 1996, *J. Leukocyte Biol.*, 59, 804-812), multiple sclerosis and brain trauma (Berman et al, 1996, *J. Immunol.*, 156, 3017-3023). An MCP-1 inhibitor may also be useful to treat stroke, reperfusion injury, ischemia, myocardial infarction and transplant rejection.

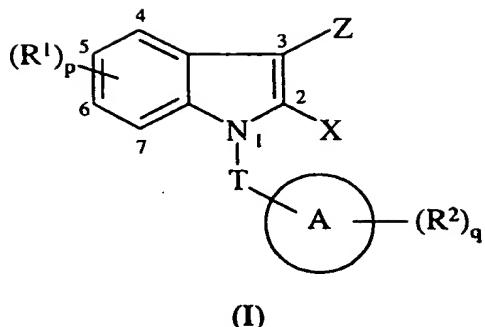
20 MCP-1 acts through the MCP-1 receptor (also known as the CCR2 receptor). MCP-2 and MCP-3 may also act, at least in part, through the MCP-1 receptor. Therefore in this specification, when reference is made to "inhibition or antagonism of MCP-1" or "MCP-1 mediated effects" this includes inhibition or antagonism of MCP-2 and/or MCP-3 mediated effects when MCP-2 and/or MCP-3 are acting through the MCP-1 receptor.

25 Japanese patent application no. JP 04273857-A discloses indole compounds for treating hypertension with a phenyl sulphonyl moiety attached to the nitrogen of the indole ring. International patent application WO96/33171 discloses similar compounds for therapy of HIV-1 infections.

The present invention is based on the discovery of a class of compounds containing an 30 indole moiety which have useful inhibitory activity against MCP-1.

Accordingly the present invention provides a compound of the formula (I)

- 2 -



which is an inhibitor of monocyte chemoattractant protein-1 and wherein:

R^1 is independently selected from trifluoromethyl, C_{1-4} alkyl, halo, hydroxy, C_{1-4} alkoxy,

5 C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, amino, cyano, C_{1-4} alkylamino, di(C_{1-4} alkyl)amino, C_{1-4} alkanoylamino, nitro, carbamoyl, C_{1-4} alkoxycarbonyl, thiol, C_{1-4} alkylsulphanyl, C_{1-4} alkylsulphinyl, C_{1-4} alkylsulphonyl, sulphonamido, carbamoyl C_{1-4} alkyl, N -(C_{1-4} alkyl)carbamoyl C_{1-4} alkyl, N -(C_{1-4} alkyl)₂carbamoyl- C_{1-4} alkyl, hydroxy C_{1-4} alkyl, C_{1-4} alkoxy C_{1-4} alkyl, morpholino, pyrrolidinyl, carboxy C_{1-4} alkylamino, R^3 and - OR^3 (where R^3

10 is optionally substituted phenyl or an optionally substituted 5- or 6- membered heteroaryl ring);

p is 1-4 and R^1 can have the same or different values when p is 2-4;

T is of the formula

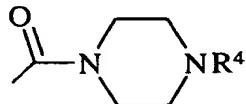


15 (where R^4 is hydrogen or C_{1-4} alkyl, m = 0-2, s = 0-2, $m + s$ = 0-2, and R^4 can take different values when $m + s$ = 2);

X is carboxy, tetrazol-5-yl, cyano, SO_3H , - SO_2NHR^4 (where R^4 is as defined above),

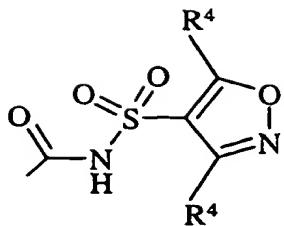
- SO_2NHAr (where Ar is an optionally substituted phenyl or optionally substituted 5 or 6 membered heteroaryl ring), - $CONHR^5$ (where R^5 is cyano, OH, - SO_2-C_{1-4} alkyl, - SO_2CF_3 ,

20 - SO_2 -phenyl, -(CHR^4)_r-COOH, (where r is 1-3 and R^4 (as defined above) can take different values when r is 2-3)), or X is a group of formula (II)



(II)

or X represents a group of formula (III)



(III)

where the groups defined as R⁴ here may have different values within the definition of R⁴ above;

A is selected from phenyl, naphthyl, furyl, pyridyl and thienyl;

5 R² is independently selected from trifluoromethyl, C₁₋₄alkyl, halo, hydroxy, trifluoromethoxy, cyano, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, amino, C₁₋₄alkylamino, di(C₁₋₄alkyl)amino, C₁₋₄alkanoylamino, nitro, carboxy, carbamoyl, C₁₋₄alkoxycarbonyl, thiol, C₁₋₄alkylsulphanyl, C₁₋₄alkylsulphanyl, C₁₋₄alkylsulphonyl, sulphonamido, carbamoylC₁₋₄alkyl, N-(C₁₋₄alkyl)carbamoylC₁₋₄alkyl, N-(C₁₋₄alkyl)₂carbamoyl-C₁₋₄alkyl, hydroxyC₁₋₄alkyl,

10 C₁₋₄alkoxyC₁₋₄alkyl or two R² values together may form a divalent radical of the formula -O(CH₂)₁₋₄O- attached to adjacent carbon atoms on ring A;

q is 0-4 and R² can have the same or different values when q is 2-4;

Z is hydrogen, fluoro, chloro, bromo, iodo, methyl, trifluoromethyl, hydroxymethyl, methoxy, methylsulphanyl, methylsulphanyl, methylsulphonyl or carboxyC₃₋₆cycloalkyl,

15 -(CHR⁴)_r-NR⁶R⁷ (where r is 0-2, R⁶ and R⁷ are independently selected from H and C₁₋₄alkyl or R⁶ and R⁷ together with the nitrogen to which they are attached form a 5 or 6 membered non-aromatic ring optionally containing one further heteroatom selected from O, N or S); or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof.

In this specification the term 'alkyl' includes straight chained, branched structures and

20 ring systems. For example, "C₁₋₄alkyl" includes propyl, isopropyl, *t*-butyl and cyclopropane. However, references to individual alkyl groups such as 'propyl' are specific for the straight chained version only, references to individual branched chain alkyl groups such as 'isopropyl' are specific for the branched chain version only and references to the cyclo groups such as cyclopropane are specific to the cyclic groups only. A similar convention applies to other 25 radicals, for example "hydroxyC₁₋₄alkyl" includes 1-hydroxyethyl and 2-hydroxyethyl. The term "halo" refers to fluoro, chloro, bromo and iodo.

Suitable optional substituents for aryl and heteroaryl are any of the values defined for R¹ and R² above. "Aryl" means phenyl or naphthyl. "Heteroaryl" means an aromatic mono- or bicyclic- 5-10 membered ring with up to three or five ring heteroatoms (in mono or bicyclic rings respectively) selected from nitrogen, oxygen and sulphur. Examples of "heteroaryl" 5 include thienyl, pyrrolyl, furanyl, imidazolyl, thiazolyl, pyrimidinyl, pyridinyl, indolyl, benzimidazolyl, benzthiazolyl, quinolyl and isoquinolinyl.

An example of "C₁₋₄alkanoyloxy" is acetoxy. Examples of "C₁₋₄alkoxycarbonyl" include methoxycarbonyl, ethoxycarbonyl, *n*- and *t*-butoxycarbonyl. Examples of "C₁₋₄alkoxy" include methoxy, ethoxy and propoxy. Examples of "C₁₋₄alkanoylamino" 10 include formamido, acetamido and propionylamino. Examples of "C₁₋₄alkylsulphanyl" include methylthio and ethylthio. Examples of "C₁₋₄alkylsulphinyl" include methylsulphinyl and ethylsulphinyl. Examples of "C₁₋₄alkylsulphonyl" include methylsulphonyl and ethylsulphonyl. Examples of "C₁₋₄alkanoyl" include propanoyl and ethanoyl. Examples of "C₁₋₄alkylamino" include methylamino and ethylamino. Examples of "di(C₁₋₄alkyl)amino" 15 include di-*N*-methylamino, di-(*N*-ethyl)amino and *N*-ethyl-*N*-methylamino. Examples of "C₁₋₄alkoxyC₁₋₄alkyl" methoxymethyl and propoxyethyl. Examples of "carbamoylC₁₋₄alkyl" are methylcarboxamide and ethylcarboxamide. Examples of "carboxyC₃₋₆cycloalkyl" are 2-carboxycyclopropyl and 3-carboxycyclopentyl. Examples of "N-(C₁₋₄alkyl)carbamoylC₁₋₄alkyl" are methylaminocarbonylethyl and 20 ethylaminocarbonylpropyl. Examples of "N-(C₁₋₄alkyl)₂carbamoyl-C₁₋₄alkyl" are dimethylaminocarbonylethyl and methylethylaminocarbonylpropyl. Examples of "carboxyC₁₋₄alkylamino" are carboxymethylamino and carboxypropylamino.

Preferred values for R¹, p, Z, X, T, A, R² and q are as follows.

Preferred values for R¹ are C₁₋₄alkoxy, halo, nitro, amino, trifluoromethyl and 25 carboxyC₁₋₄alkylamino, more preferably chloro and/or C₁₋₄alkoxy. Where R¹ is halo, fluoro, chloro or bromo are preferred. Where R¹ is C₁₋₄alkoxy it is preferably methoxy or ethoxy, particularly methoxy. Preferably position 7 is unsubstituted, and preferably there is no more than one C₁₋₄alkoxy group.

Preferably p is 1 or 2.

30 Preferred combinations of p and R¹ are as follows.

When p=1 then R¹ is preferably fluoro, chloro or methoxy and in particular 5-chloro and 6-chloro.

T is preferably $-\text{SO}_2^-$.

Preferably X is carboxy, $-\text{CONHR}^5$ (where R^5 is $-\text{SO}_2-\text{C}_{1-4}\text{alkyl}$, $-\text{SO}_2\text{CF}_3$, $-\text{SO}_2\text{-phenyl}$) or tetrazol-5-yl. R^5 is preferably $-\text{SO}_2\text{CF}_3$. In particular X is carboxy.

Preferably A is phenyl, naphthyl, furyl and thienyl in particular phenyl or thienyl.

5 When A is thienyl it is preferably thien-2-yl. Most preferably A is phenyl.

R^2 is preferably chloro, bromo, methyl, methoxy, nitro, trifluoromethyl or trifluoromethoxy. Another preferred value for R^2 is fluoro.

q is preferably 1 or 2, especially 2.

Preferred combinations of A, R^2 and q are as follows.

10 When A is phenyl, and q is 1, then R^2 is preferably chloro especially 3-chloro or 4-chloro. Other preferred values for R^2 include 3-fluoro, 4-fluoro and 3-trifluoromethyl.

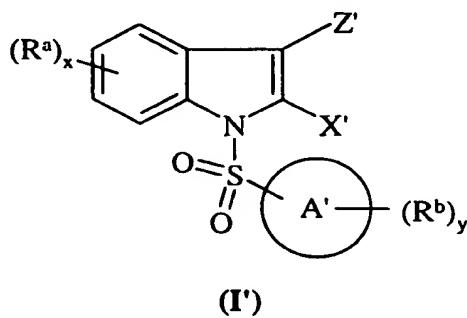
When A is phenyl, and q is 2, then R^2 is preferably chloro, especially 3,4-dichlorophenyl. Another preferred value is fluoro, especially 3,4-difluoro.

When A is phenyl then the positions ortho to T are preferably unsubstituted.

15 When A is thien-2-yl then preferably R^2 is chloro, especially 5-chloro.

Preferably Z is hydrogen or bromo, especially hydrogen.

Therefore a preferred class of compounds is that of formula (I'):



20 wherein;

R^a is methoxy, fluoro, chloro, bromo, nitro, amino, trifluoromethyl or carboxymethylamino;

x is 1 or 2 with the proviso that there is at most one methoxy group;

X' is carboxy, $-\text{CONHSO}_2\text{CF}_3$ or tetrazol-5-yl;

25 A' is phenyl or thienyl;

R^b is chloro, bromo, methyl, methoxy, nitro, trifluoromethyl or trifluoromethoxy;

y is 1 or 2;

Z' is hydrogen or bromo;

or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof.

Preferably R^a is chloro or methoxy. Preferably position 7 is unsubstituted.

Preferred combinations of x and R^a are as follows.

5 When x=1 then R^a is preferably chloro or methoxy especially 5-chloro or 6-chloro.

Z' is preferably hydrogen.

A' is preferably phenyl. Where A' is thienyl it is preferably thien-2-yl.

Preferred combinations of A', R^b and y are as follows.

When A' is phenyl, and y is 1, then R^b is preferably chloro in particular 3-chloro or

10 4-chloro. Another preferred value is fluoro, in particular 3-fluoro or 4-fluoro.

When A' is phenyl, and y is 2, then R^b is preferably chloro, in particular

3,4-dichlorophenyl. Another preferred value is fluoro, in particular 3,4-difluoro.

When A' is phenyl then the positions ortho to the SO₂ moiety linked to the indole ring are preferably hydrogen.

15 When A' is thien-2-yl then preferably R^b is chloro in particular 5-chloro.

Preferred compounds having formula (I) or (IA) (defined below) include any one of:

N-(3,4-Dichlorophenylsulphonyl)-5-chloroindole-2-carboxylic acid;

N-(6-Bromonaphthalen-2-ylsulphonyl)-5-chloroindole-2-carboxylic acid;

N-(3-Chlorophenylsulphonyl)-5-chloroindole-2-carboxylic acid; and

20 3-Bromo-5-Fluoro-*N*-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid;

or an *in vivo* hydrolysable ester or a pharmaceutically acceptable salt thereof.

Suitable pharmaceutically acceptable salts include acid addition salts such as methanesulfonate, fumarate, hydrochloride, hydrobromide, citrate, maleate and salts formed with phosphoric and sulphuric acid. In another aspect suitable salts are base salts such as an alkali metal salt for example sodium, an alkaline earth metal salt for example calcium or magnesium, an organic amine salt for example triethylamine, morpholine, *N*-methylpiperidine, *N*-ethylpiperidine, procaine, dibenzylamine, *N,N*-dibenzylethylamine or amino acids for example lysine. There may be more than one cation or anion depending on the number of charged functions and the valency of the cations or anions. A preferred 25 pharmaceutically acceptable salt is a sodium salt.

Some compounds of formula (I) may possess chiral centres. It is to be understood that the invention encompasses all such optical isomers and diastereoisomers of compounds of formula (I).

The invention further relates to all tautomeric forms of the compounds of formula (I).

5 It is also to be understood that certain compounds of the formula (I) can exist in solvated as well as unsolvated forms such as, for example, hydrated forms. It is to be understood that the invention encompasses all such solvated forms.

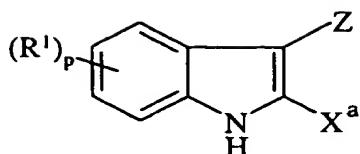
An *in vivo* hydrolysable ester of a compound of the formula (I) containing carboxy or hydroxy group is, for example, a pharmaceutically acceptable ester which is hydrolysed in the 10 human or animal body to produce the parent acid or alcohol.

Suitable pharmaceutically acceptable esters for carboxy include C₁₋₆alkoxymethyl esters for example methoxymethyl, C₁₋₆alkanoyloxymethyl esters for example pivaloyloxymethyl, phthalidyl esters, C₃₋₈cycloalkoxy-carbonyloxyC₁₋₆alkyl esters for example 1-cyclohexylcarbonyloxyethyl; 1,3-dioxolen-2-onylmethyl esters for example 15 5-methyl-1,3-dioxolen-2-onylmethyl; and C₁₋₆alkoxycarbonyloxyethyl esters for example 1-methoxycarbonyloxyethyl and may be formed at any carboxy group in the compounds of this invention.

An *in vivo* hydrolysable ester of a compound of the formula (I) containing a hydroxy group includes inorganic esters such as phosphate esters and α -acyloxyalkyl ethers and related 20 compounds which as a result of the *in vivo* hydrolysis of the ester breakdown to give the parent hydroxy group. Examples of α -acyloxyalkyl ethers include acetoxymethoxy and 2,2-dimethylpropionyloxymethoxy. A selection of *in vivo* hydrolysable ester forming groups for hydroxy include alkanoyl, benzoyl, phenylacetyl and substituted benzoyl and phenylacetyl, alkoxycarbonyl (to give alkyl carbonate esters), dialkylcarbamoyl and 25 N-(dialkylaminoethyl)-N-alkylcarbamoyl (to give carbamates), dialkylaminoacetyl and carboxyacetyl.

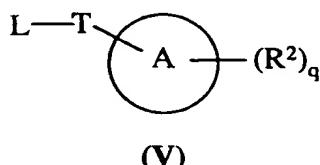
Another aspect of the present invention provides a process for preparing a compound of formula (I) or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof which process comprises of:

30 a) reacting compounds of formula (IV):



(IV)

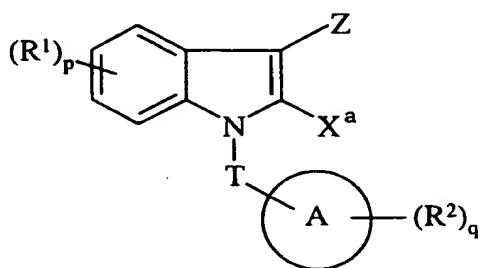
where X^a is carboxy protected in the form of an ester and other groups are as defined in formula (I) with a compound of formula (V):



5

(V)

where L is a leaving group other groups are as defined in formula (I) to give a compound of formula (VI):



10

(VI)

where X^a is carboxy protected as an ester; and

b) optionally interconverting a compound of formula (VI) to give another compound of formula (VI), wherein any functional groups are protected if necessary and optionally:

- removing any protecting groups;
- optionally forming a pharmaceutically acceptable salt or *in vivo* hydrolysable ester.

Compounds of formula (VI) and (I) may be interconverted for example as described herein or by known processes such as functional group modification or aromatic substitution.

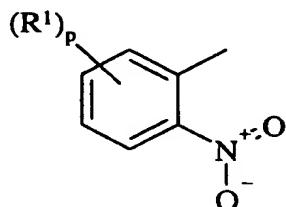
L is a leaving group. Preferable values for L are chloro and bromo.

Compounds of formula (IV) and (V) may be reacted together in an inert solvent and a base such as *N,N*-dimethylformamide/sodium hydride or dichloromethane/sodium hydroxide (optionally in the presence of a phase transfer catalyst such as tetra-*n*-butylammonium hydrogensulphate) for 1-6 hours preferably 1-3 hours, at a temperature of 15-30°C, preferably 20-25°C to give a compound of formula (VI).

Compounds of formula (IV) are commercially available, made by modification using known processes of commercially available compounds of formula (IV), or they are prepared by:

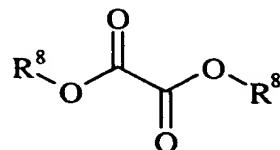
a) Reacting a compound of formula (VII):

5



(VII)

where R^1 and p are as defined in formula (I), with a compound of formula (VIII)

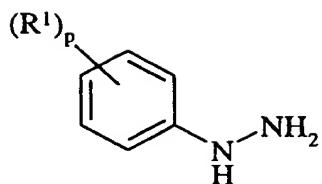


10 (VIII)

where R^8 is C_{1-4} alkyl.

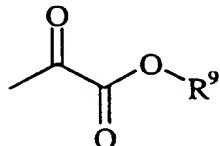
Compounds of formula (VII) and (VIII) are reacted together under Reissert reaction conditions such as in an inert solvent (such as tetrahydrofuran), in the presence of a base (such as potassium ethoxide), at a temperature range of 15-30°C preferably 20-25°C, for 10-20 15 hours preferably 15-17 hours. The resulting compound is isolated and dissolved in an alcohol such as ethanol and an organic acid (such as acetic acid) and a transition metal catalyst (such as 10% Pd/C) and cyclohexene is added. The mixture is heated at a temperature of 60-120°C preferably at 70-90°C for 15-25 hours preferably 16-20 hours to give a compound of formula (VI) in which Z is hydrogen. Then if desired Z can be optionally converted into another value 20 of Z as defined in formula (I) using techniques known in the art such as those described below.

b) Reacting a compound of formula (IX):



(IX)

where R^1 and p are as defined for formula (I), with a compound of formula (X):



(X)

5 where R^9 is C_{1-4} alkyl.

Compounds of formula (IX) and (X) are reacted together under Fischer conditions

such as with an organic acid (such as acetic acid), in an alcohol (such as ethanol), at a

temperature of $60-90^\circ C$ preferably $75-85^\circ C$, for 1-5 hours preferably 1-3 hours. The resulting

10 compound is mixed with a strong acid (such as polyphosphoric acid) and heated at $90-150^\circ C$ preferably $100-120^\circ C$ for 0.5-4 hours preferably 0.5-2 hours to give a compound of formula (VI) in which Z is hydrogen. Then if desired Z can be optionally converted into another value of Z as defined in formula (I) using techniques known in the art such as those described below.

15 Compounds of formula (V), (VII), (VIII), (IX) and (X) are known or commercially available or are prepared by processes known in the art by standard manipulation of commercially available or known materials.

R^8 and R^9 are C_{1-4} alkyl. Preferably R^8 and R^9 are methyl or ethyl.

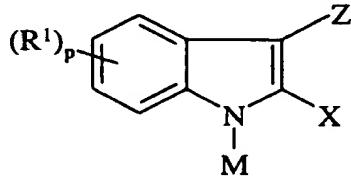
It will be appreciated that certain of the various optional substituents in the compounds

20 of the present invention may be introduced by standard aromatic substitution reactions or generated by conventional functional group modifications either prior to or immediately following the processes mentioned above, and as such are included in the process aspect of the invention. Such reactions and modifications include, for example, introduction of a substituent by means of an aromatic substitution reaction, reduction of substituents, alkylation 25 of substituents and oxidation of substituents. The reagents and reaction conditions for such procedures are well known in the chemical art. Particular examples of aromatic substitution

reactions include the introduction of a nitro group using concentrated nitric acid, the introduction of an acyl group using, for example, an acyl halide and Lewis acid (such as aluminium trichloride) under Friedel Crafts conditions; the introduction of an alkyl group using an alkyl halide and Lewis acid (such as aluminium trichloride) under Friedel Crafts conditions; and the introduction of a halogeno group. Particular examples of modifications include the reduction of a nitro group to an amino group by for example, catalytic hydrogenation with a nickel catalyst or treatment with iron in the presence of hydrochloric acid with heating; oxidation of alkylthio to alkylsulphinyl or alkylsulphonyl. Specific examples of the substitution and modification reactions prior to or immediately following the processes mentioned above are illustrated, but not limited by, the following examples in which variable groups are as defined for formula (I) unless otherwise stated.

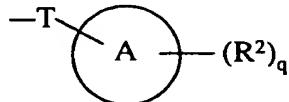
1) Modification of R¹.

a) For R¹ = Ar: compounds of formula (XI)

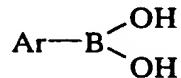


15 (XI)

M is H, a nitrogen protecting group or the group



where R¹ is Br are coupled with compounds of formula (XII)



20 (XII)

where Ar is an optionally substituted phenyl or optionally substituted 5 or 6 membered heteroaryl ring, to give compounds of formula (XI) where R¹ = Ar. Suitable reaction conditions are set out below.

Compounds of formula (XI) where R¹ = Br and (XII) are reacted together in the

25 presence of a transition metal catalyst (for example tetrakis(triphenylphosphine)palladium(0)), in an inert solvent (such as toluene) and an alcohol (such as ethanol), with an aqueous base

(such as potassium carbonate), preferably in an inert atmosphere, at a temperature of 60-100°C preferably 75-85°C for 14-20 hours preferably 15-17 hours.

b) For $R^1 = NH_2$; compounds of formula (XI) where $R^1 = NO_2$ are reduced under standard conditions to give a compound of formula (XI) where $R^1 = NH_2$. Suitable reaction conditions 5 are set out below.

Compounds of formula (XI) where $R^1 = NO_2$ are reacted with a reducing agent (such as sodium borohydride) and stannous chloride dihydrate in an alcohol (such as ethanol) at a temperature of 30-80°C preferably 50-70°C for 2-10 hours preferably 4-6 hours.

c) For $R^1 = MeC(O)NH-$: compounds of formula (XI) where $R^1 = MeC(O)NH-$ can be 10 prepared from compounds of formula (XI) where $R^1 = NH_2$. Suitable reaction conditions are set out below.

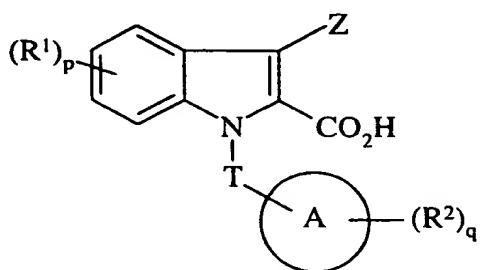
Compounds of formula (XI) where $R^1 = NH_2$ are reacted in acetic anhydride at a temperature of 60-140°C preferably 80-100°C for 0.5-5 hours preferably 0.5-2 hours.

d) For $R^1 = C_{1-4}alkoxycarbonylC_{1-4}alkylamino$: compounds of formula (XI) where 15 $R^1 = C_{1-4}alkoxycarbonylC_{1-4}alkylamino$ can be prepared from compounds of formula (XI) where $R^1 = NH_2$. Suitable reaction conditions are set out below.

Compounds of formula (XI) where $R^1 = NH_2$ are reacted with the corresponding glyoxalate, aldehyde ester or keto ester (such as ethyl glyoxalate) followed by the addition of a reducing agent (such as sodium cyanoborohydride) in an alcohol (such as ethanol) with an 20 acid (such as acetic acid) for 1-10 minutes preferably 4-6 minutes at 15-30°C preferably 20-25°C.

2) Modification of X.

a) For X = carboxy: hydrolysing a compound of formula (VI) as defined above to give a compound of formula (XIII):



Suitable reaction conditions are set out below.

Compounds of formula (VI) where X^a is $-\text{CO}_2\text{Me}$ may be conveniently hydrolysed to give compounds of formula (XIII) with a salt (such as lithium iodide), in an organic base (such as pyridine), at a temperature range of 100-125°C, in particular 115-120°C, for 3-10

5 hours, preferably 5-7 hours followed by the addition of aqueous acid (for example 2M hydrochloric acid).

b) For $X = -\text{CONHR}^5$ (R^5 is as defined for formula (I)): compounds where $X = -\text{CONHR}^5$ can be prepared by coupling compounds of formula (XIII) and compounds of formula (XIV):



10 (XIV)

under standard peptide coupling conditions. Suitable reaction conditions are set out below.

Compounds of formula (XIII) and compounds of formula (XIV) can be reacted together in the presence of a coupling agent (such as

1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride) and optionally a catalyst

15 (such as dimethylamino pyridine) in an inert solvent (such as dichloromethane) for 1-36 hours preferably 20-30 hours at 15-30°C preferably 20-25°C.

c) For $X = \text{C(O)NH}_2$ and CN: compounds of formula (XI) where $X = \text{C(O)NH}_2$ and CN can be prepared from compounds of formula (XI) where $X = \text{carboxy}$. Suitable reaction conditions are set out below.

20 Compounds of formula (XI) where $X = \text{carboxy}$ are reacted with ammonia in an organic base (such as pyridine) with a sulphonating agent (such as methanesulphonyl chloride) at a temperature of -10 to 10°C preferably -2 to 2°C for 1-5 hours preferably 2-3 hours to give both compounds.

d) For $X = \text{tetrazol-5-yl}$: compounds of formula (XI) where $X = \text{tetrazol-5-yl}$ can be prepared 25 from compounds of formula (XI) where $X = \text{CN}$. Suitable reaction conditions are set out below.

Compounds of formula (XI) where $X = \text{CN}$ are reacted with an azide (such as sodium azide) and triethylamine hydrochloride in a solvent (such as *N*-methyl pyrrolidinone) at a temperature of 100-200°C preferably 140-160°C for 3-10 hours preferably 5-6 hours.

30 3) Modification of Z.

a) For Z = Br: compounds of formula (XI) where Z = hydrogen may be brominated under standard conditions to give a compound of formula (XI) where Z = Br. Suitable reaction conditions are set out below.

Compounds of formula (XI) where Z = bromine may be prepared by reacting a 5 compound of formula (XI) where Z = hydrogen in an inert solvent (such as N,N-dimethylformamide) with bromine for 5-55 minutes particularly 25-35 minutes at 10-30°C, preferably 20-25°C.

Other modifications of Z are achieved by standard reactions known in the art. For example:

10 a) a substituent of the formula -CH₂NR₂ can be prepared by the Mannich reaction.

Compounds of formula (XI) where Z = hydrogen are treated with formaldehyde and an amine in the presence of acid;

b) the Vilsmeier formulation of compounds of formula (XI) where Z = hydrogen with POCl₃ and N,N-dimethylformamide gives the aldehyde in the three position which can then be 15 selectively reduced to the carbinol (with NaBH₄) or to the methyl (with NaBH₄ and trifluoroacetic acid) under standard reaction conditions.

The reader is also directed to Japanese patent application no. JP 04273857-A and International patent application WO 96/33171 for synthetic details of sulphonyl indole compounds.

20 It will also be appreciated that in some of the reactions mentioned herein it may be necessary/desirable to protect any sensitive groups in the compounds. The instances where protection is necessary or desirable and suitable methods for protection are known to those skilled in the art. Thus, if reactants include groups such as amino, carboxy or hydroxy it may be desirable to protect the group in some of the reactions mentioned herein.

25 A suitable protecting group for an amino or alkylamino group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an alkoxy carbonyl group, for example a methoxycarbonyl, ethoxycarbonyl or *t*-butoxycarbonyl group, an arylmethoxycarbonyl group, for example benzyloxycarbonyl, or an aroyl group, for example benzoyl. The deprotection conditions for the above protecting groups necessarily vary with the choice of protecting

30 group. Thus, for example, an acyl group such as an alkanoyl or alkoxy carbonyl group or an aroyl group may be removed for example, by hydrolysis with a suitable base such as an alkali metal hydroxide, for example lithium or sodium hydroxide. Alternatively an acyl group such

as a *t*-butoxycarbonyl group may be removed, for example, by treatment with a suitable acid as hydrochloric, sulphuric or phosphoric acid or trifluoroacetic acid and an arylmethoxycarbonyl group such as a benzyloxycarbonyl group may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon, or by treatment with a Lewis acid for example boron tris(trifluoroacetate). A suitable alternative protecting group for a primary amino group is, for example, a phthaloyl group which may be removed by treatment with an alkylamine, for example dimethylaminopropylamine, or with hydrazine.

A suitable protecting group for a hydroxy group is, for example, an acyl group, for example an alkanoyl group such as acetyl, an aroyl group, for example benzoyl, or an arylmethyl group, for example benzyl. The deprotection conditions for the above protecting groups will necessarily vary with the choice of protecting group. Thus, for example, an acyl group such as an alkanoyl or an aroyl group may be removed, for example, by hydrolysis with a suitable base such as an alkali metal hydroxide, for example lithium or sodium hydroxide. Alternatively an arylmethyl group such as a benzyl group may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

A suitable protecting group for a carboxy group is, for example, an esterifying group, for example a methyl or an ethyl group which may be removed, for example, by hydrolysis with a base such as sodium hydroxide, or for example a *t*-butyl group which may be removed, for example, by treatment with an acid, for example an organic acid such as trifluoroacetic acid, or for example a benzyl group which may be removed, for example, by hydrogenation over a catalyst such as palladium-on-carbon.

The protecting groups may be removed at any convenient stage in the synthesis using conventional techniques well known in the chemical art.

When a pharmaceutically-acceptable salt of a compound of the formula (I) is required, it may be obtained, for example, by reaction of said compound with the appropriate acid (which affords a physiologically acceptable anion), or with the appropriate base (which affords a physiologically acceptable cation), or by any other conventional salt formation procedure.

When an optically active form of a compound of the formula (I) is required, it may be obtained, for example, by carrying out one of the aforesaid procedures using an optically active starting material or by resolution of a racemic form of said compound using a conventional procedure.

According to a further aspect of the invention there is provided a compound of the formula (I) as defined herein, or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof, for use in a method of treatment of the human or animal body by therapy.

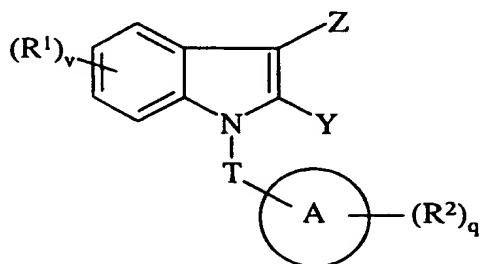
The invention also provides a compound of formula (I) as defined herein, or a

5 pharmaceutically acceptable salt, or an *in vivo* hydrolysable ester thereof, for use as a medicament.

Another aspect of the present invention provides the use of a compound of formula (I) or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof, in the manufacture of a medicament for use in antagonising an MCP-1 mediated effect in a warm

10 blooded animal, such as man.

Another aspect of the present invention provides the use of a compound of formula (IA)



15

(IA)

where R¹, Z, T, A, R² and q are as defined for formula (I);

Y = X (as defined for formula (I)) or -CONHR¹⁰ (where R¹⁰ is H or C₁₋₄alkyl);

v = 0-4;

a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof, in the manufacture
20 of a medicament for use in antagonising an MCP-1 mediated effect in a warm blooded animal, such as man.

A further aspect of the present invention comprises a novel compound of the formula (IA) or a pharmaceutically acceptable salt thereof or an *in vivo* hydrolysable ester thereof, for example a compound of the formula (IA) as defined above in which Y is carboxy, T is -SO₂-,
25 A(R²)_q is phenyl independently substituted at the 3 and 4 positions by halogeno (such as 3,4-dichlorophenyl or 3,4-difluorophenyl), v is 1 or 2 and R¹ is attached at the 4 and/or 5 position of the indole ring. Particular novel compounds of formula (IA) include, for example,

N-(3-chlorophenylsulphonyl)indole-2-carboxylic acid;
N-(3,4-Dichlorophenylsulphonyl)indole-2-carboxylic acid;
N-(4,5-Dichlorothien-2-ylsulphonyl)indole-2-carboxylic acid;
3-Bromo-*N*-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid; and

5 3-Chloro-*N*-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid;
or a pharmaceutically acceptable salt thereof or an *in vivo* hydrolysable ester thereof.

According to a further aspect of the present invention there is provided a method for antagonising an MCP-1 mediated effect in a warm blooded animal, such as man, in need of such treatment, which comprises administering to said animal an effective amount of a

10 compound of formula (IA), or a pharmaceutically acceptable salt, or an *in vivo* hydrolysable ester thereof. According to a further aspect of the invention there is provided a method of inhibiting the binding of MCP-1 to a receptor thereof in a warm blooded animal in need thereof which comprises administering to said animal an effective amount of a compound of formula (I) or (IA), or a pharmaceutically acceptable salt, or an *in vivo* hydrolysable ester

15 thereof. According to a further aspect of the invention there is provided the use of a compound of formula (I) or (IA) for the manufacture of a medicament for use in inhibiting the binding of MCP-1 to a receptor thereof.

In order to use a compound of the formula (I) (or (IA)) or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof for the therapeutic treatment of

20 mammals including humans, in particular in treating inflammation, it is normally formulated in accordance with standard pharmaceutical practice as a pharmaceutical composition.

Therefore in another aspect the present invention provides a pharmaceutical composition which comprises a compound of the formula (I) (or novel compound of formula (IA)) or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof and a

25 pharmaceutically acceptable diluent or carrier.

The compositions of the invention may be in a form suitable for oral use (for example as tablets, lozenges, hard or soft capsules, aqueous or oily suspensions, emulsions, dispersible powders or granules, syrups or elixirs), for topical use (for example as creams, ointments, gels, or aqueous or oily solutions or suspensions), for administration by inhalation (for

30 example as a finely divided powder or a liquid aerosol), for administration by insufflation (for example as a finely divided powder) or for parenteral administration (for example as a sterile

aqueous or oily solution for intravenous, subcutaneous, intramuscular or intramuscular dosing or as a suppository for rectal dosing).

The compositions of the invention may be obtained by conventional procedures using conventional pharmaceutical excipients, well known in the art. Thus, compositions intended 5 for oral use may contain, for example, one or more colouring, sweetening, flavouring and/or preservative agents.

Suitable pharmaceutically acceptable excipients for a tablet formulation include, for example, inert diluents such as lactose, sodium carbonate, calcium phosphate or calcium carbonate, granulating and disintegrating agents such as corn starch or algenic acid; binding 10 agents such as starch; lubricating agents such as magnesium stearate, stearic acid or talc; preservative agents such as ethyl or propyl p-hydroxybenzoate, and anti-oxidants, such as ascorbic acid. Tablet formulations may be uncoated or coated either to modify their disintegration and the subsequent absorption of the active ingredient within the 15 gastrointestinal track, or to improve their stability and/or appearance, in either case, using conventional coating agents and procedures well known in the art.

Compositions for oral use may be in the form of hard gelatin capsules in which the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules in which the active ingredient is mixed with water or an oil such as peanut oil, liquid paraffin, or olive oil.

20 Aqueous suspensions generally contain the active ingredient in finely powdered form together with one or more suspending agents, such as sodium carboxymethylcellulose, methylcellulose, hydroxypropylmethylcellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents such as lecithin or condensation products of an alkylene oxide with fatty acids (for example polyoxyethylene stearate), or 25 condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters 30 derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions

may also contain one or more preservatives (such as ethyl or propyl *p*-hydroxybenzoate, anti-oxidants (such as ascorbic acid), colouring agents, flavouring agents, and/or sweetening agents (such as sucrose, saccharine or aspartame).

Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil (such as arachis oil, olive oil, sesame oil or coconut oil) or in a mineral oil (such as liquid paraffin). The oily suspensions may also contain a thickening agent such as beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set out above, and flavouring agents may be added to provide a palatable oral preparation. These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

10 Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water generally contain the active ingredient together with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above. Additional excipients such as sweetening, flavouring and colouring agents, may also be present.

15 The pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, such as olive oil or arachis oil, or a mineral oil, such as for example liquid paraffin or a mixture of any of these. Suitable emulsifying agents may be, for example, naturally-occurring gums such as gum acacia or gum tragacanth, naturally-occurring phosphatides such as soya bean, lecithin, an esters or partial 20 esters derived from fatty acids and hexitol anhydrides (for example sorbitan monooleate) and condensation products of the said partial esters with ethylene oxide such as polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening, flavouring and preservative agents.

Syrups and elixirs may be formulated with sweetening agents such as glycerol, 25 propylene glycol, sorbitol, aspartame or sucrose, and may also contain a demulcent, preservative, flavouring and/or colouring agent.

The pharmaceutical compositions may also be in the form of a sterile injectable aqueous or oily suspension, which may be formulated according to known procedures using one or more of the appropriate dispersing or wetting agents and suspending agents, which 30 have been mentioned above. A sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example a solution in 1,3-butanediol.

Suppository formulations may be prepared by mixing the active ingredient with a suitable non-irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Suitable excipients include, for example, cocoa butter and polyethylene glycols.

5 Topical formulations, such as creams, ointments, gels and aqueous or oily solutions or suspensions, may generally be obtained by formulating an active ingredient with a conventional, topically acceptable, vehicle or diluent using conventional procedure well known in the art.

Compositions for administration by insufflation may be in the form of a finely divided 10 powder containing particles of average diameter of, for example, 30 μ or much less, the powder itself comprising either active ingredient alone or diluted with one or more physiologically acceptable carriers such as lactose. The powder for insufflation is then conveniently retained in a capsule containing, for example, 1 to 50mg of active ingredient for use with a turbo-inhaler device, such as is used for insufflation of the known agent sodium 15 cromoglycate.

Compositions for administration by inhalation may be in the form of a conventional 20 pressurised aerosol arranged to dispense the active ingredient either as an aerosol containing finely divided solid or liquid droplets. Conventional aerosol propellants such as volatile fluorinated hydrocarbons or hydrocarbons may be used and the aerosol device is conveniently arranged to dispense a metered quantity of active ingredient.

For further information on Formulation the reader is referred to Chapter 25.2 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The amount of active ingredient that is combined with one or more excipients to 25 produce a single dosage form will necessarily vary depending upon the host treated and the particular route of administration. For example, a formulation intended for oral administration to humans will generally contain, for example, from 0.5 mg to 2 g of active agent compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight of the total composition. Dosage unit forms will 30 generally contain about 1 mg to about 500 mg of an active ingredient. For further information on Routes of Administration and Dosage Regimes the reader is referred to Chapter 25.3 in

Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The size of the dose for therapeutic or prophylactic purposes of a compound of the Formula I will naturally vary according to the nature and severity of the conditions, the age 5 and sex of the animal or patient and the route of administration, according to well known principles of medicine. As mentioned above, compounds of the Formula I are useful in treating diseases or medical conditions which are due alone or in part to the effects of farnesylation of rats.

In using a compound of the Formula I for therapeutic or prophylactic purposes it will 10 generally be administered so that a daily dose in the range, for example, 0.5 mg to 75 mg per kg body weight is received, given if required in divided doses. In general lower doses will be administered when a parenteral route is employed. Thus, for example, for intravenous administration, a dose in the range, for example, 0.5 mg to 30 mg per kg body weight will generally be used. Similarly, for administration by inhalation, a dose in the range, for 15 example, 0.5 mg to 25 mg per kg body weight will be used. Oral administration is however preferred.

The following illustrate, but are not intended to limit, representative pharmaceutical dosage forms of the invention as defined herein (the active ingredient being termed "Compound X"), for therapeutic or prophylactic use in humans:

20 (a)

<u>Tablet I</u>	<u>mg/tablet</u>
Compound X.	100
Lactose Ph.Eur	182.75
Croscarmellose sodium	12.0
Maize starch paste (5% w/v paste)	2.25
Magnesium stearate	3.0

(b)

<u>Tablet II</u>	<u>mg/tablet</u>
Compound X	50
Lactose Ph.Eur	223.75
Croscarmellose sodium	6.0
Maize starch	15.0
Polyvinylpyrrolidone (5% w/v paste)	2.25
Magnesium stearate	3.0

(c)

<u>Tablet III</u>	<u>mg/tablet</u>
Compound X	1.0
Lactose Ph.Eur	93.25
Croscarmellose sodium	4.0
Maize starch paste (5% w/v paste)	0.75
Magnesium stearate	1.0

5

(d)

<u>Capsule</u>	<u>mg/capsule</u>
Compound X	10
Lactose Ph.Eur	488.5
Magnesium	1.5

(e)

<u>Injection I</u>	<u>(50 mg/ml)</u>
Compound X	5.0% w/v
1M Sodium hydroxide solution	15.0% v/v
0.1M Hydrochloric acid	to adjust pH to 7.6
Polyethylene glycol 400	4.5% w/v
Water for injection	to 100%

(f)

<u>Injection II</u>	(10 mg/ml)
Compound X	1.0% w/v
Sodium phosphate BP	3.6% w/v
0.1M Sodium hydroxide solution	15.0% v/v
Water for injection	to 100%

(g)

<u>Injection III</u>	(1mg/ml, buffered to pH6)
Compound X	0.1% w/v
Sodium phosphate BP	2.26% w/v
Citric acid	0.38% w/v
Polyethylene glycol 400	3.5% w/v
Water for injection	to 100%

5

(h)

<u>Aerosol I</u>	<u>mg/ml</u>
Compound X	10.0
Sorbitan trioleate	13.5
Trichlorofluoromethane	910.0
Dichlorodifluoromethane	490.0

(i)

<u>Aerosol II</u>	<u>mg/ml</u>
Compound X	0.2
Sorbitan trioleate	0.27
Trichlorofluoromethane	70.0
Dichlorodifluoromethane	280.0
Dichlorotetrafluoroethane	1094.0

(j)

<u>Aerosol III</u>	<u>mg/ml</u>
Compound X	2.5
Sorbitan trioleate	3.38
Trichlorofluoromethane	67.5
Dichlorodifluoromethane	1086.0
Dichlorotetrafluoroethane	191.6

(k)

<u>Aerosol IV</u>	<u>mg/ml</u>
Compound X	2.5
Soya lecithin	2.7
Trichlorofluoromethane	67.5
Dichlorodifluoromethane	1086.0
Dichlorotetrafluoroethane	191.6

5

(l)

<u>Ointment</u>	<u>ml</u>
Compound X	40 mg
Ethanol	300 µl
Water	300 µl
1-Dodecylazacycloheptan-2-one	50 µl
Propylene glycol	to 1 ml

Note

The above formulations may be obtained by conventional procedures well known in the pharmaceutical art. The tablets (a)-(c) may be enteric coated by conventional means, for example to provide a coating of cellulose acetate phthalate. The aerosol formulations (h)-(k) may be used in conjunction with standard, metered dose aerosol dispensers, and the suspending agents sorbitan trioleate and soya lecithin may be replaced by an alternative

suspending agent such as sorbitan monooleate, sorbitan sesquioleate, polysorbate 80, polyglycerol oleate or oleic acid.

Biological Testing.

Abbreviations:

ATCC	American Type Culture Collection, Rockville, USA.
BCA	bicinchoninic acid, (used, with copper sulphate, to assay protein)
DMEM	Dulbecco's modified Eagle's medium
EGTA	ethylenebis(oxyethylenenitrilo)tetraacetic acid
FCS	foetal calf serum
HBSS	Hank's Balanced Salt Solution
hMCP-1	human Monocyte Chemoattractant Protein-1
PBS	phosphate buffered saline
PCR	polymerase chain reaction

5 AMPLITAQ™, available from Perkin-Elmer Cetus, is used as the source of thermostable DNA polymerase.

Binding Buffer is 50 mM HEPES, 1 mM CaCl₂, 5 mM MgCl₂, 0.5% foetal calf serum, adjusted to pH 7.2 with 1 M NaOH.

Non-Essential Amino Acids (100X concentrate) is: L-Alanine, 890 mg/l;

10 L-Asparagine, 1320 mg/l; L-Aspartic acid, 1330 mg/l; L-Glutamic acid, 1470 mg/l; Glycine, 750 mg/l; L-Proline, 1150 mg/l and; L-Serine, 1050 mg/l.

Hypoxanthine and Thymidine Supplement (50x concentrate) is: hypoxanthine, 680 mg/l and; thymidine, 194 mg/l.

15 Penicillin-Streptomycin is: Penicillin G (sodium salt); 5000 units/ml; Streptomycin sulphate, 5000 µg/ml.

Human monocytic cell line THP-1 cells are available from ATCC, accession number ATCC TIB-202.

20 Hank's Balanced Salt Solution (HBSS) was obtained from Gibco; see *Proc. Soc. Exp. Biol. Med.*, 1949, 71, 196.

Synthetic cell culture medium, RPMI 1640 was obtained from Gibco; it contains inorganic salts [Ca(NO₃)₂.4H₂O 100 mg/l; KCl 400 mg/l; MgSO₄.7H₂O 100 mg/l; NaCl 6000 mg/l; NaHCO₃ 2000 mg/l & Na₂HPO₄ (anhyd) 800 mg/l], D-Glucose 2000 mg/l, reduced glutathione 1 mg/l, amino acids and vitamins.

FURA-2/AM is

1-[2-(5-carboxyoxazol-2-yl)-6-aminobenzofuran-5-oxy]-2-(2'-amino-5'-methylphenoxy)-ethane-N,N,N',N'-tetraacetic acid pentaacetoxymethyl ester and was obtained from Molecular Probes, Eugene, Oregon, USA.

5 General molecular biology procedures can be followed from any of the methods described in "Molecular Cloning-A Laboratory Manual" Second Edition, Sambrook, Fritsch and Maniatis (Cold Spring Harbor Laboratory, 1989).

Biological Assays for hMCP-1 Antagonists

a) hMCP-1 Receptor-binding assay

10 **i) Cloning and expression of hMCP-1 receptor**

The MCP-1 receptor B (CCR2B) cDNA was cloned by PCR from THP-1 cell RNA using suitable oligonucleotide primers based on the published MCP-1 receptor sequences (Charo *et al.*, 1994, *Proc. Natl. Acad. Sci. USA*, **91**, 2752). The resulting PCR products were cloned into vector PCR-II™ (InVitrogen, San Diego, CA.). Error free CCR2B cDNA was 15 subcloned as a Hind III-Not I fragment into the eukaryotic expression vector pCDNA3 (InVitrogen) to generate pCDNA3/CC-CKR2A and pCDNA3/CCR2B respectively.

Linearised pCDNA3/CCR2B DNA was transfected into CHO-K1 cells by calcium phosphate precipitation (Wigler *et al.*, 1979, *Cell*, **16**, 777). Transfected cells were selected by the addition of Geneticin Sulphate (G418, Gibco BRL) at 1mg/ml, 24 hours after the cells had 20 been transfected. Preparation of RNA and Northern blotting were carried out as described previously (Needham *et al.*, 1995, *Prot. Express. Purific.*, **6**, 134). CHO-K1 clone 7 (CHO-CCR2B) was identified as the highest MCP-1 receptor B expressor.

ii) Preparation of membrane fragments

CHO-CCR2B cells were grown in DMEM supplemented with 10% foetal calf serum, 25 2 mM glutamine, 1x Non-Essential Amino Acids, 1x Hypoxanthine and Thymidine Supplement and Penicillin-Streptomycin (at 50 µg streptomycin/ml, Gibco BRL). Membrane fragments were prepared using cell lysis/differential centrifugation methods as described previously (Siciliano *et al.*, 1990, *J. Biol. Chem.*, **265**, 19658). Protein concentration was estimated by BCA protein assay (Pierce, Rockford, Illinois) according to the manufacturer's 30 instructions.

iii) Assay

125I MCP-1 was prepared using Bolton and Hunter conjugation (Bolton *et al.*, 1973, *Biochem. J.*, **133**, 529; Amersham International plc]. Equilibrium binding assays were carried out using the method of Ernst *et al.*, 1994, *J. Immunol.*, **152**, 3541. Briefly, varying amounts of ¹²⁵I-labeled MCP-1 were added to 10 mg of purified CHO-CCR2B cell membranes in 100

5 ml of Binding Buffer. After 1 hour incubation at room temperature the binding reaction mixtures were filtered and washed 5 times through a plate washer (Packard Harvester Filtermate™ 196). Scintillation fluid (25µl, Microscint™-20, a high efficiency liquid scintillation counting cocktail for aqueous samples) was added to each well and the plate was covered with plate sealer and counted (Packard Top Count™). Cold competition studies were

10 performed as above using 100 pM ¹²⁵I-labeled MCP-1 in the presence of varying concentrations of unlabelled MCP-1. Non-specific binding was determined by the inclusion of a 200-fold molar excess of unlabelled MCP-1 in the reaction.

Ligand binding studies with membrane fragments prepared from CHO-CCR2B cells showed that the CCR2B was present at a concentration of 0.2 pmoles/mg of membrane

15 protein and bound MCP-1 selectively and with high affinity ($IC_{50} = 110$ pM, $K_d = 120$ pM). Binding to these membranes was completely reversible and reached equilibrium after 45 minutes at room temperature, and there was a linear relationship between MCP-1 binding and CHO-CCR2B cell membrane concentration when using MCP-1 at concentrations between 100 pM and 500 pM.

20 Test compounds dissolved in DMSO (5 µl) were tested in competition with 100 pM labelled MCP-1 over a concentration range (0.1-200µM) in duplicate using eight point dose-response curves and IC_{50} concentrations were calculated.

b) MCP-1 mediated calcium flux in THP-1 cells

The human monocytic cell line THP-1 was grown in a synthetic cell culture medium

25 RPMI 1640 supplemented with 10 % foetal calf serum, 2 mM glutamine and Penicillin-Streptomycin (at 50 µg streptomycin/ml, Gibco BRL). THP-1 cells were washed in HBSS (lacking Ca^{2+} and Mg^{2+}) + 1 mg/ml BSA and resuspended in the same buffer at a density of 3×10^6 cells/ml. The cells were then loaded with 1 mM FURA-2/AM for 30 min at 37°C, washed twice in HBSS, and resuspended at 1×10^6 cells/ml. THP-1 cell suspension (0.9

30 ml) was added to a 5 ml disposable cuvette containing a magnetic stirrer bar and 2.1 ml of prewarmed (37°C) HBSS containing 1 mg/ml BSA, 1 mM $MgCl_2$ and 2 mM $CaCl_2$. The

cuvette was placed in a fluorescence spectrophotometer (Perkin Elmer, Norwalk, CT) and preincubated for 4 min at 37°C with stirring. Fluorescence was recorded over 70 sec and cells were stimulated by addition of hMCP-1 to the cuvette after 10 sec. $[Ca^{2+}]_i$ was measured by excitation at 340 nm and 380 nm alternately and subsequent measurement of the intensity of 5 the fluorescence emission at 510 nm. The ratio of the intensities of the emitted fluorescent light following excitation at 340 nm and 380 nm, (R), was calculated and displayed to give and estimate of cytoplasmic $[Ca^{2+}]$ according to the equation:-

$$[Ca^{2+}]_i = K_d \frac{(R - R_{min})}{(R_{max} - R)} (Sf2/Sb2)$$

10 where the K_d for FURA-2 Ca^{2+} complex at 37°C was taken to be 224 nm. R_{max} is the maximal fluorescence ratio determined after addition of 10 mM Ionomycin, R_{min} is the minimal ratio determined by the subsequent addition of a Ca^{2+} free solution containing 5 mM EGTA, and Sf2/Sb2 is the ratio of fluorescence values at 380 nm excitation determined at R_{min} and R_{max} , respectively.

15 Stimulation of THP-1 cells with hMCP-1 induced a rapid, transient rise in $[Ca^{2+}]_i$ in a specific and dose dependent manner. Dose response curves indicated an approximate EC_{50} of 2 nm. Test compounds dissolved in DMSO (10 μ l) were assayed for inhibition of calcium release by adding them to the cell suspension 10 sec prior to ligand addition and measuring the reduction in the transient rise in $[Ca^{2+}]_i$. Test compounds were also checked for lack of 20 agonism by addition in place of hMCP-1.

c) **hMCP-1 mediated chemotaxis assay.**

In vitro chemotaxis assays were performed using either the human monocytic cell line THP-1 or peripheral blood mixed monocytes obtained from fresh human blood purified by erythrocyte sedimentation followed by density gradient centrifugation over 9.6 % (w/v) sodium 25 metrizoate and 5.6 % (w/v) polysaccharide, density 1.077 g/ml (Lymphoprep™ Nycomed). Cell migration through polycarbonate membranes was measured by enumerating those passing through either directly by Coulter counting or indirectly by use of a colourimetric viability assay measuring the cleavage of a tetrazolium salt by the mitochondrial respiratory chain (Scudiero D.A. *et al.* 1988, *Cancer Res.*, **48**, 4827-4833).

30 Chemoattractants were introduced into a 96-well microtiter plate which forms the lower well of a chemotaxis chamber fitted with a PVP-free 5 μ m poresize polycarbonate

adhesive framed filter membrane (NeuroProbe MB series, Cabin John, MD 20818, USA) according to the manufacturer's instructions. The chemoattractant was diluted as appropriate in synthetic cell culture medium, RPMI 1640 (Gibco) supplemented with 2 mM glutamine and 0.5% BSA. Each dilution was degassed under vacuum for 30 min and was placed (400 µl) 5 in the lower wells of the chamber and THP-1 cells (5×10^5 in 100 µl RPMI 1640 + 0.5% BSA) were incubated in each well of the upper chamber. For the inhibition of chemotaxis the chemoattractant was kept at a constant submaximal concentration determined previously for each chemokine and added to the lower well together with the test compounds dissolved in DMSO (final DMSO concentration < 0.05% v/v) at varying concentrations. The chamber was 10 incubated for 2 h at 37°C under 5 % CO₂. The medium was removed from the upper wells which were then washed out with 200 µl physiological saline before opening the chamber, wiping dry the membrane surface and centrifuging the 96-well plate at 600 g for 5 min to harvest the cells. Supernatant (150 µl) was aspirated and 10 µl of cell proliferation reagent, WST-1, {4-[3-(4-iodophenyl)-2-(4-nitrophenyl)-2H-5-tetrazolio]-1,3-phenyl disulfonate} plus 15 an electron coupling reagent (Boehringer Mannheim, Cat.no. 1644 807) was added back to the wells. The plate was incubated at 37°C for 3 h and the absorbance of the soluble formazan product was read on a microtitre plate reader at 450 nm. The data was input into a spreadsheet, corrected for any random migration in the absence of chemoattractant and the average absorbance values, standard error of the mean, and significance tests were calculated. 20 hMCP-1 induced concentration dependent cell migration with a characteristic biphasic response, maximal 0.5-1.0 nm.

Compounds tested of the present invention generally had IC₅₀ values of less than 50µM in the hMCP-1 receptor binding assay described herein. For example the compound of example 2.01 had an IC₅₀ of 10µM.

25 The invention is further illustrated, but not limited by the following Examples.

General Procedures.

N,N-Dimethylformamide (DMF) was dried over 4Å molecular sieves. Anhydrous tetrahydrofuran (THF) was obtained from Aldrich SURESEAL™ bottles. Other commercially available reagents and solvents were used without further purification unless otherwise stated. 30 Organic solvent extracts were dried over anhydrous MgSO₄. ¹H, ¹³C and ¹⁹F NMR were recorded on Bruker WM200, WM250, WM300 or WM400 instruments using Me₂SO-δ₆ with Me₄Si or CCl₃F as internal standard as appropriate, unless otherwise stated. Chemical shifts

are in δ (ppm) and peak multiplicities are designated as follows: s, singlet; d, doublet; dd, doublet of doublets; t, triplet; dt, doublet of triplets; q, quartet; m, multiplet; br, broad. Mass spectra were recorded on VG 12-12 quadrupole, VG 70-250 SE, VG ZAB 2-SE or a VG modified AEI/Kratos MS9 spectrometers. For TLC analysis, Merck precoated TLC plates 5 (silica gel 60 F254, d = 0.25 mm) were used. Flash chromatography was performed on silica (Merck Kieselgel: Art.9385). Melting point determinations were performed on a Kofler block or with a Büchi melting point apparatus and are uncorrected. All temperatures are in degrees centigrade.

Example 1

10 Methyl N-(3,4-dichlorophenylsulphonyl)indole-2-carboxylate

Methyl indole-2-carboxylate (0.15 g) was dissolved in DMF and sodium hydride (41 mg) was added in a single portion. The reaction was stirred for 1 hour, then 3,4-dichlorobenzenesulphonyl chloride (0.25 g) was added in a single portion. Stirring was continued for a further 2 hours and then the reaction was quenched by the addition of water. 15 The reaction mixture was partitioned between water and ethyl acetate. Combined organic extracts were dried (MgSO_4) and concentrated *in vacuo* and the residue purified by column chromatography using isohexane-5% ethyl acetate as eluent to give the desired end product as a solid (51%). NMR δ (CDCl_3) 3.96 (s, 3H), 7.22-8.20 (m, 8H); M/z (+) 384 ($M\text{H}^+$), 352, 175.

Examples 1.01 - 1.05.

20 The procedure described in Example 1 was repeated using the appropriate indole-2-carboxylic ester and arylsulphonyl halide. Thus there were obtained the compounds described below.

Example 1.01: Methyl N-(3-chlorophenylsulphonyl)indole-2-carboxylate in 82% yield; NMR δ (CDCl_3) 3.91 (s, 3H), 7.20-8.14 (m, 9H); M/z (+) 350 ($M\text{H}^+$), 318, 175, 144.

Example 1.02: Methyl N-(4,5-dichlorothien-2-ylsulphonyl)indole-2-carboxylate in 19% yield;

25 NMR δ (CDCl_3) 3.98 (s, 3H), 7.22-8.08 (m, 6H); M/z (+) 390 ($M\text{H}^+$), 358, 175, 144.

Example 1.03: Methyl 5-chloro-N-(3,4-dichlorophenylsulphonyl)-indole-2-carboxylate in 24% yield; NMR δ (CDCl_3) 3.87 (s, 3H), 7.44 (s, 1H), 7.54 (dd, 1H), 7.82 (d, 1H), 7.96 (m, 2H), 8.12 (d, 1H), 8.28 (d, 1H).

Example 1.04: Methyl N-(6-bromonaphthalen-2-ylsulphonyl)-5-chloroindole-2-carboxylate in

30 9% yield; NMR δ (CDCl_3) 3.88 (s, 3H), 7.39 (s, 1H), 7.52 (dd, 1H), 7.78 (d, 1H), 7.84 (dd, 1H), 7.94 (dd, 1H), 8.12 (m, 2H), 8.20 (d, 1H), 8.35 (s, 1H), 8.82 (s, 1H).

Example 1.05: Methyl N-(3-chlorophenylsulphonyl)-5-chloroindole-2-carboxylate in 24% yield; NMR δ (CDCl₃) 3.87 (s, 3H), 7.41 (s, 1H), 7.53 (dd, 1H), 7.68 (t, 1H), 7.82 (m, 2H), 7.98 (d, 1H), 8.04 (s, 1H), 8.09 (d, 1H).

5 **Example 1.06:** Methyl 3-bromo-N-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylate in 31% yield; NMR δ (CD₃SOCD₃) 3.96 (s, 3H), 7.46 (t, 1H), 7.57 (dd, 1H), 7.62 (d, 1H), 7.88 (t, 1H), 8.1 (d, 1H), 8.14 (d, 1H), 8.21 (s, 1H), 8.26 (d, 1H).

Example 1.07: Methyl 4-acetoxy-N-(3,4-dichlorophenylsulphonyl)indole-2-carboxylate in 65% yield; NMR δ (CD₃SOCD₃) 2.34 (s, 3H), 3.86 (s, 3H), 7.16 (d, 1H), 7.50 (s, 1H), 7.55 (t, 1H), 7.92-8.06 (m, 3H), 8.31 (d, 1H); *M/z* (+) 442 (MH⁺).

10 **Example 1.08:** Methyl 3-chloro-N-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylate in 68% yield; NMR δ (CD₃SOCD₃) 3.96 (s, 3H), 7.48 (t, 1H), 7.59-7.68 (m, 2H), 7.88 (t, 1H), 8.1-8.16 (m, 2H), 8.2 (s, 1H), 8.26 (d, 1H); *M/z* (+) 418 (MH⁺).

15 **Example 1.09:** Methyl 3-chloro-5-fluoro-N-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylate in 34% yield; NMR δ (CD₃SOCD₃) 3.98 (s, 3H), 7.37-7.49 (m, 2H), 7.89 (t, 1H), 8.13-8.18 (m, 2H), 8.22 (s, 1H), 8.27 (d, 1H).

Example 2

N-(3-Chlorophenylsulphonyl)indole-2-carboxylic acid.

Methyl *N*-(3-chlorophenylsulphonyl)indole-2-carboxylate (0.56 g) and lithium iodide (2.0 g) were dissolved in pyridine and heated at reflux for 6 hours, cooled to room temperature and poured into 2M HCl and extracted with diethyl ether. Combined organic extracts were dried (MgSO₄) and concentrated *in vacuo* to give an oil which was purified by column chromatography using DCM-2% methanol as eluent to give the desired product as a white solid (0.24 g, 45%), mp 216-217°; NMR δ (CD₃SOCD₃) 7.30-8.10 (m, 9H); *M/z*(-) 334 (M-H⁺), 290, 226, 191, 180, 116.

Examples 2.01-2.05.

The procedure described in Example 2 was repeated using the appropriate indole-2-carboxylic ester. Thus there were obtained the compounds described below.

30 **Example 2.01:** N-(3,4-Dichlorophenylsulphonyl)indole-2-carboxylic acid in 74% yield, mp 203-204°; NMR δ (CD₃SOCD₃) 7.30 (m, 2H), 7.5 (m, 1H), 7.7 (m, 1H), 7.9 (m, 1H), 8.0 (m, 2H), 8.25 (m, 1H); *M/z*(-) 370 (M⁺),

Example 2.02: *N*-(4,5-Dichlorothien-2-ylsulphonyl)indole-2-carboxylic acid in 75% yield, mp 82-183°; NMR δ (CD₃SOCD₃) 8.25 (s, 1H), 7.95 (d, 1H), 7.63 (d, 1H), 7.42 (t, 1H), 7.25 (t, 1H), 7.10 (m, 1H); *M/z*(-) 376 (*M*⁺), 374, 332, 330, 268, 266, 233, 231.

Example 2.03: *N*-(3,4-Dichlorophenylsulphonyl)-5-chloroindole-2-carboxylic acid in 57% yield; NMR δ (CD₃SOCD₃) 7.12 (s, 1H), 7.44 (dd, 1H), 7.72 (d, 1H), 7.88 (d, 1H), 8.02 (d, 1H), 8.08 (dd, 1H), 8.38 (s, 1H); *M/z*(-) 404 (*M*-H⁺), 358.

Example 2.04: *N*-(6-Bromonaphthalen-2-ylsulphonyl)-5-chloroindole-2-carboxylic acid in 68% yield; NMR δ (CD₃SOCD₃) 6.92 (s, 1H), 7.36 (dd, 1H), 7.64 (s, 1H), 7.80 (dd, 1H), 8.07 (m, 2H), 8.16 (m, 2H), 8.32 (s, 1H), 8.86 (s, 1H); *M/z*(-) 466 (*M*-H⁺), 464, 462, 418.

Example 2.05: *N*-(3-Chlorophenylsulphonyl)-5-chloroindole-2-carboxylic acid in 68% yield; NMR δ (CD₃SOCD₃) 7.26 (s, 1H), 7.48 (dd, 1H), 7.66 (t, 1H), 7.77 (s, 1H), 7.82 (d, 1H), 8.05 (t, 2H), 8.12 (s, 1H); *M/z*(-) 368 (*M*-H⁺), 324.

Example 2.06: 3-Bromo-*N*-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid in 50% yield; NMR δ (CD₃SOCD₃) 7.42 (t, 1H), 7.5-7.59 (m, 2H), 7.88 (t, 1H), 8.08 (d, 1H), 8.12 (d, 1H), 8.32-8.38 (m, 2H); *M/z*(-) 446 (*M*-H⁺), 402, 322.

Example 2.07: 3-Chloro-*N*-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid in 58% yield; NMR δ (CD₃SOCD₃) 7.45 (s, 1H), 7.56 (d, 1H), 7.62 (d, 1H), 7.88 (t, 1H), 8.11 (t, 2H), 8.28-8.36 (m, 2H); *M/z*(-) 402 (*M*-H⁺), 360, 358.

Example 2.08: 3-Bromo-5-Fluoro-*N*-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid in 69% yield; NMR δ (CD₃SOCD₃) 7.32-7.46 (m, 2H), 7.88 (t, 1H), 8.08-8.16 (m, 2H), 8.31-8.37 (m, 2H); *M/z*(-) 466 (*M*-H⁺), 404, 422, 420.

Example 3

Methyl *N*-(3,4-dichlorophenylsulphonyl)-4-hydroxyindole-2-carboxylate.

Aqueous sodium hydrogencarbonate solution (16 ml, 50%) was added to methyl 4-acetoxy-*N*-(3,4-dichlorophenylsulphonyl)indole-2-carboxylate (0.63 g) in methanol (15 ml) and the reaction stirred for 48 hours at room temperature. The solution was then poured into 2M HCl and extracted with ethyl acetate. Combined organic extracts were dried (MgSO₄) and concentrated *in vacuo* to give the desired product as a gum (0.5 g, 87%); NMR δ (CD₃SOCD₃) 3.83 (s, 3H), 6.71 (d, 1H), 7.31 (t, 1H), 7.42-7.5 (m, 2H), 7.92 (s, 2H), 8.19 (s, 1H), 10.31 (s, 1H); *M/z*(+) 402 (*M*H⁺), 400.

Preparation of Starting Materials.

Starting materials for the Examples above are either commercially available or are readily prepared by standard methods from known materials, for example the following reactions (Methods A to F) are illustrations but not limitations of the preparation of some of the starting materials used in the above reactions.

Method A**6-Bromo-2-naphthylsulphonyl chloride**

A solution of sodium nitrite (2.7 g) in water (5 ml) was added during 2 hours to a stirred mixture of 6-amino-2-naphthalene-sulphonic acid (8.8 g), dilute aqueous hydrochloric acid (2.8% weight/volume, 20 ml) and water (15 ml) which had been cooled to 0°C. The mixture was stirred at 0°C for 30 minutes and poured onto a stirred suspension of cuprous bromide (5.34 g) in dilute aqueous hydrobromic acid (2.8%, 20 ml). The mixture was stored at ambient temperature for 18 hours. The mixture was evaporated to give 6-bromo-2-naphthalenesulphonic acid which was used without further purification.

The material was suspended in DMF (40 ml) and cooled to 5°C. Thionyl chloride (8.6 ml) was added dropwise and the mixture was stirred at 5°C for 3 hours. The mixture was poured onto ice and extracted with methylene chloride. The organic solution was dried ($MgSO_4$) and evaporated. The residue was purified by column chromatography using a 20:1 mixture of hexane and ethyl acetate as eluent to give the desired starting material in 22% yield. NMR δ (CD_3SOCD_3) 7.65 (m, 1H), 7.75-8.0 (m, 3H), 8.15-8.2 (m, 2H).

Method B**Ethyl 3-bromoindole-2-carboxylate**

A solution of bromine (2.72 ml) in DMF was added dropwise over 10 mins to a solution of ethyl indole-2-carboxylate in DMF. The reaction was stirred for 30 mins, then poured into water to precipitate a pale yellow solid which was filtered off and recrystallized from ethyl acetate to give the desired starting material as white needles (10.2 g, 72%), mp 150-151°C; NMR δ ($CDCl_3$) 1.44 (t, 3H), 4.45 (q, 2H), 7.22 (m, 1H), 7.38 (m, 2H), 7.66 (d, 1H), 9.27 (bs, 1H); M/z (-) 268 (M^+), 266, 196, 194.

The procedure described above was repeated using the appropriate indole. Thus was obtained the compound described below.

Methyl 3-bromo-5-fluoroindole-2-carboxylate in 83% yield; NMR δ (CD_3SOCD_3) 3.9 (s, 3H), 7.08-7.28 (m, 2H), 7.49 (dd, 1H), 12.38 (bs, 1H); M/z (+) 274 (MH^+), 272.

Method C**Ethyl 3-chloroindole-2-carboxylate**

Ethyl 3-chloroindole-2-carboxylate (3 g) and phosphorous pentachloride (9 g) were heated at 90 °C for 1 hour. The mixture was then cooled to room temperature, poured into 5 water and the resulting solid filtered purified by column chromatography using isohexane-20% ethyl acetate as eluent to give the desired end product as a white solid (1.25 g, 35%); NMR δ (CD₃SOCD₃) 1.3 (t, 3H), 4.4 (q, 2H), 7.2 (t, 1H), 7.35 (t, 1H), 7.45 (d, 1H), 7.6 (d, 1H), 12.5 (1H, bs); *M/z* (-) 222 (*M-H*⁺).

Method D**10 Methyl-5-chloroindole-2-carboxylate**

Sodium (10.3 g, 447 mmol) was dissolved in methanol (HPLC grade, anhydrous, 150 ml) with continual stirring under an atmosphere of argon. Upon complete dissolution of the sodium ethyl-5-chloroindole-2-carboxylate (10.08 g, 44.7 mmol) was added as a single portion and the reaction mixture was stirred at room temperature for 2 hours. The reaction 15 mixture was acidified with the addition of aqueous hydrochloric acid (excess) causing the precipitation of a white solid. The solid was filtered and washed with aqueous hydrochloric acid (100 ml) and water (100mls) then dried overnight in at 55° to yield the product as a white solid (8.97 g, 95%). NMR δ (CD₃SOCD₃) 3.86 (s, 3H), 7.12 (dd, 1H), 7.24 (dd, 1H), 7.43 (d, 1H), 7.72 (s, 1H), 12.10 (brs, 1H).

20 The procedure described above was repeated using the appropriate indole. Thus was obtained the compound described below.

Methyl 3-bromoindole-2-carboxylate in 79% yield; NMR δ (CD₃SOCD₃) 3.90 (s, 3H), 7.18 (t, 1H), 7.35 (t, 1H), 7.47 (d, 1H), 7.53 (d, 1H), 12.24 (brs, 1H).

Methyl 3-chloroindole-2-carboxylate in 64% yield; NMR δ (CD₃SOCD₃) 3.90 (s, 3H), 7.18 (t, 1H), 7.35 (t, 1H), 7.45 (d, 1H), 7.59 (d, 1H); *M/z* (+) 212 (*MH*⁺), 210.

Method E**Methyl 4-hydroxyindole-2-carboxylate**

Boron tribromide (73.1 ml, 1.0 M solution in DCM) was added dropwise to a solution of methyl 4-methoxyindole-2-carboxylate (5 g) in DCM (200 ml) cooled to -78°C under 30 argon. The reaction was allowed to warm to room temperature then partitioned between dichloromethane and saturated aqueous sodium hydrogen carbonate solution. Combined

organic extracts were dried ($MgSO_4$) and concentrated *in vacuo* and the residue purified by column chromatography using isohexane-50% ethyl acetate as eluent to give the end product as a yellow solid (2.98 g, 64%); NMR δ (CD_3SOCD_3) 3.82 (s, 3H), 6.36 (d, 1H), 6.85 (d, 1H), 7.02 (t, 1H), 7.17 (d, 1H), 9.66 (s, 1H), 11.72 (bs, 1H); M/z (+) 192 (MH^+).

5 **Method F**

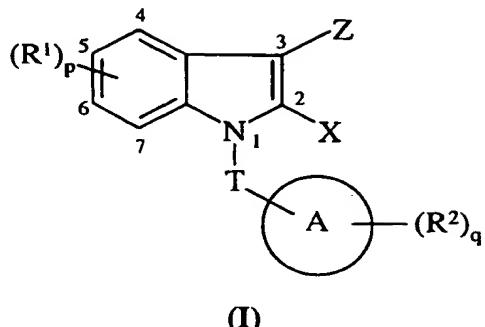
Methyl 4-acetoxyindole-2-carboxylate

Methyl 4-hydroxyindole-2-carboxylate (0.5 g) and 4-dimethylaminopyridine (50 mg) were dissolved in acetic anhydride (5 ml) and heated at 80^0C for 3 hours. The reaction was allowed to cool overnight to precipitate white crystals, which were filtered and dried *in vacuo* (0.44 g, 72%); NMR δ (CD_3SOCD_3) 2.34 (s, 3H), 3.85 (s, 3H), 6.80 (d, 1H), 7.06 (s, 1H), 7.23 (t, 1H), 7.29-7.35 (m, 1H), 12.1 (bs, 1H); M/z (-) 232 ($M-H^+$).

CLAIMS:

What we claim is:

5 1. A compound of the formula (I)



wherein

10 R^1 is independently selected from trifluoromethyl, C_{1-4} alkyl, halo, hydroxy, C_{1-4} alkoxy, C_{1-4} alkanoyl, C_{1-4} alkanoyloxy, amino, cyano, C_{1-4} alkylamino, di(C_{1-4} alkyl)amino, C_{1-4} alkanoylamino, nitro, carbamoyl, C_{1-4} alkoxycarbonyl, thiol, C_{1-4} alkylsulphanyl, C_{1-4} alkylsulphanyl, C_{1-4} alkylsulphonyl, sulphonamido, carbamoyl C_{1-4} alkyl, $N-(C_{1-4}\text{alkyl})\text{carbamoyl}C_{1-4}\text{alkyl}$, $N-(C_{1-4}\text{alkyl})_2\text{carbamoyl-}C_{1-4}\text{alkyl}$, hydroxy C_{1-4} alkyl, C_{1-4} alkoxy C_{1-4} alkyl, morpholino, pyrrolidinyl, carboxy C_{1-4} alkylamino, R^3 and - OR^3 (where R^3 is optionally substituted phenyl or an optionally substituted 5- or 6- membered heteroaryl ring);

15 p is 1-4 and R^1 can have the same or different values when p is 2-4;

T is of the formula

20 $-(\text{CHR}^4)_m-(\text{SO}_2)-(\text{CHR}^4)_s-$,
(where R^4 is hydrogen or C_{1-4} alkyl, m = 0-2, s = 0-2, $m + s$ = 0-2, and R^4 can take different values when $m + s$ = 2);

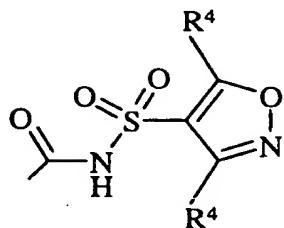
X is carboxy, tetrazol-5-yl, cyano, SO_3H , $-\text{SO}_2\text{NHR}^4$ (where R^4 is as defined above), $-\text{SO}_2\text{NAr}$ (where Ar is an optionally substituted phenyl or optionally substituted 5 or 6 membered heteroaryl ring), $-\text{CONHR}^5$ (where R^5 is cyano, OH, $-\text{SO}_2\text{-}C_{1-4}\text{alkyl}$, $-\text{SO}_2\text{CF}_3$, $-\text{SO}_2\text{-phenyl}$, $-(\text{CHR}^4)_r\text{-COOH}$, (where r is 1-3 and R^4 (as defined above) can take different values when r is 2-3)), or X is a group of formula (II)



(II)

or X represents a group of formula (III)

5



(III)

where the groups defined as R⁴ here may have different values within the definition of R⁴ above;

A is selected from phenyl, naphthyl, furyl, pyridyl and thienyl;

10 R² is independently selected from trifluoromethyl, C₁₋₄alkyl, halo, hydroxy, trifluoromethoxy, cyano, C₁₋₄alkoxy, C₁₋₄alkanoyl, C₁₋₄alkanoyloxy, amino, C₁₋₄alkylamino, di(C₁₋₄alkyl)amino, C₁₋₄alkanoylamino, nitro, carboxy, carbamoyl, C₁₋₄alkoxycarbonyl, thiol, C₁₋₄alkylsulphanyl, C₁₋₄alkylsulphinyl, C₁₋₄alkylsulphonyl, sulphonamido, carbamoylC₁₋₄alkyl, N-(C₁₋₄alkyl)carbamoylC₁₋₄alkyl, N-(C₁₋₄alkyl)₂carbamoyl-C₁₋₄alkyl, hydroxyC₁₋₄alkyl,

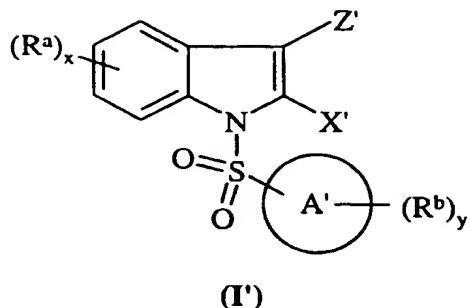
15 C₁₋₄alkoxyC₁₋₄alkyl or two R² values together may form a divalent radical of the formula -O(CH₂)₁₋₄O- attached to adjacent carbon atoms on ring A;

q is 0-4 and R² can have the same or different values when q is 2-4;

Z is hydrogen, fluoro, chloro, bromo, iodo, methyl, trifluoromethyl, hydroxymethyl, methoxy, methylsulphanyl, methylsulphinyl, methylsulphonyl or carboxyC₃₋₆cycloalkyl,

20 -(CHR⁴)_r-NR⁶R⁷ (where r is 0-2, R⁶ and R⁷ are independently selected from H and C₁₋₄alkyl or R⁶ and R⁷ together with the nitrogen to which they are attached form a 5 or 6 membered non-aromatic ring optionally containing one further heteroatom selected from O, N or S); or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof.

25 2. A compound as claimed in claim 1 which is a compound of the formula (I').



wherein

R^a is methoxy, fluoro, chloro, bromo, nitro, amino, trifluoromethyl or carboxymethylamino;

5 x is 1 or 2 with the proviso that there is at most one methoxy group;

X' is carboxy, $-\text{CONHSO}_2\text{CF}_3$ or tetrazol-5-yl;

A' is phenyl or thienyl;

R^b is chloro, bromo, methyl, methoxy, nitro, trifluoromethyl or trifluoromethoxy;

y is 1 or 2;

10 Z' is hydrogen or bromo;

or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof.

3. A compound as claimed in claim 1 or 2 wherein $A(R^2)_q$ or $A'(R^b)_y$ is 3-chlorophenyl, 4-chlorophenyl, 3-fluorophenyl, 4-fluorophenyl, 3-trifluoromethylphenyl, 3,4-dichlorophenyl 15 or 3,4-difluorophenyl.

4. A compound as claimed in claim 1, 2 or 3 wherein X or X' is carboxy.

5. A compound as claimed in claim 1 which is selected from

20 $N-(3,4\text{-dichlorophenylsulphonyl})-5\text{-chloroindole-2-carboxylic acid};$

$N-(6\text{-bromonaphthalene-2-ylsulphonyl})-5\text{-chloroindole-2-carboxylic acid};$

$N-(3\text{-chlorophenylsulphonyl})-5\text{-chloroindole-2-carboxylic acid};$ and

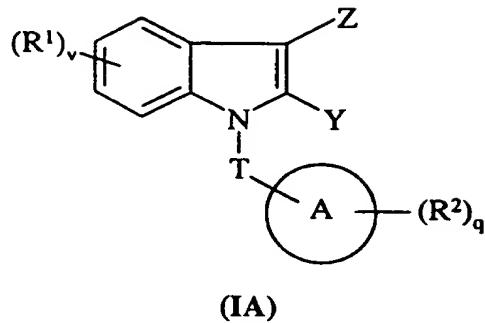
$3\text{-bromo-5-fluoro-N-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid};$

or an *in vivo* hydrolysable ester or a pharmaceutically acceptable salt thereof.

25

6. A pharmaceutical composition which comprises a compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, together with a pharmaceutically acceptable diluent or carrier.

7. The use of a compound of the formula (IA).



wherein

5 R^1 , Z , T , A , R^2 and q have any of the values defined in claim 1;
 Y has any of the values defined for X in claim 1 or Y is $-CONHR^{10}$ where R^{10} is H or
 C_{1-4} alkyl; and
 $v = 0-4$;
or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof, in the
10 manufacture of a medicament for use in antagonising an MCP-1 mediated effect in a warm
blooded animal.

8. A compound of the formula (IA) as defined in claim 7, or a pharmaceutically
acceptable salt thereof, wherein Y is carboxy; T is $-SO_2-$; $A(R^2)_q$ is phenyl independently
15 substituted at the 3 and 4 positions by halogeno; v is 1 or 2; and R^1 is attached at the 4 and/or
5 position of the indole ring.

9. A compound of the formula (IA) as defined in claim 7 which is selected from
N-(3-chlorophenylsulphonyl)indole-2-carboxylic acid;
20 N-(3,4-dichlorophenylsulphonyl)indole-2-carboxylic acid;
N-(4,5-dichlorothien-2-ylsulphonyl)indole-2-carboxylic acid;
3-bromo-N-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid; and
3-chloro-N-(3-trifluoromethylphenylsulphonyl)indole-2-carboxylic acid; or a
pharmaceutically acceptable salt thereof or an *in vivo* hydrolysable ester thereof.

10. A pharmaceutical composition which comprises a compound as claimed in claim 8 or 9, or a pharmaceutically acceptable salt thereof or *in vivo* hydrolysable ester thereof, together with a pharmaceutically acceptable diluent or carrier.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 98/02340

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 C07D209/42 A61K31/40 C07D409/12

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 IPC 6 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	DE 39 43 225 A (SCHERING AG) 27 June 1991 see page 5, line 1 ---	1
X	YUUSAKU YOKOYAMA T AL.: "Palladium-catalyzed cross-coupling reaction: direct allylation of aryl bromides with allyl acetate" TETRAHEDRON LETTERS., vol. 26, no. 52, - 1985 pages 6457-6460, XP002081581 OXFORD GB * page 6458-6459: compound 7 * --- -/-	1

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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- "&" document member of the same patent family

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Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.
Fax: (+31-70) 340-3016

Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

...International Application No.

PCT/GB 98/02340

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>CHEMICAL ABSTRACTS, vol. 123, no. 14, 2 October 1995 Columbus, Ohio, US; abstract no. 179521d, KATAOKA, KENICHIRO ET AL.: "Homopiperazines as cell migration inhibitors." XP002081582 see abstract & JP 95 145060 A (TEJIN LTD.) -----</p>	1,6,7

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/GB 98/02340

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			WO	9109858 A	11-07-1991
			EP	0460153 A	11-12-1991
			JP	4505928 T	15-10-1992
			PT	96351 A	30-09-1991
			US	5254563 A	19-10-1993

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